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To:

Bradley L. Sisson

Primary Patent Examiner

FIRM:

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(703) 308-4242 or (703) 305-3014

FROM:

Dan Hart

CLIENT CODE:

GENSET.020APC

PAGES:

14 (INCLUDING THIS SHEET)

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MESSAGE:

As we recently discussed, attached please find an Amendment containing the claims we discussed in our telephonic interview on March 29, 2001. I will call you to confirm your receipt of this fax.

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PATENT

Case Docket No. GENSET.020APC Date: April 10, 2001

Page 1

In re application of:

Cohen, et al.

App. No.

09/463,075

Filed

January 14, 2000

For

BIALLELIC MARKERS FOR

USE IN CONSTRUCTING A

HIGH DENSITY

DISEQUILIBRIUM MAP OF THE

HUMAN GENOME

Examiner

B. Sisson

Art Unit

1655

I hereby certify that this correspondence and all marked attachments are being facetimile transmitted to the Patent and Trademark Office on the date shown below:

April 10, 2001

Daniel Hart, Reg. No. 40,637

ASSISTANT COMMISSIONER FOR PATENT'S WASHINGTON, D.C. 20231

Sir:

Transmitted herewith is an amendment in the above-identified application.

The fee has been calculated as shown below:

CLAIMS AS FILED							
	CLAIMS REMAINING AFTER AMENDMENT		HIGHEST NO. PREVIOUSLY PAID FOR	SLY EXTRA		ADDITIONAL FEE	
Total Claims	20	_	38	= 0	×	\$18	= \$0
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If application has bee dependent claim(s),		ıtain multiple				\$270	= \$0
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Daniel Hart

Registration No. 40,637

Attorney of Record

GENSET.020APC

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Applicant	:	Cohen et al.)	Group Art Unit 1655					
Appl. No.	:	09/463,075)	CERTIFICATE OF FAX TRANSMISSION					
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For	:	BIALLELIC MARKERS FOR USE IN CONSTRUCTING A HIGH DENSITY DISEQUILIBRIUM MAP OF THE HUMAN GENOME)))	April 10, 2001 (Date) Danlet Hart, Reg. No. 40,637					
Examiner	:	B. Sisson	_	1.11					
Assistant Commissioner for Patents									

Assistant Commissioner for Patents Washington, D.C. 20231

Dear Sir:

In response to the Office Action mailed January 10, 2001 in the above-identified application, please amend the application as follows.

IN THE SPECIFICATION:

Please replace the paragraph beginning on page 17, line 5 with the following paragraph:

Alternatively, the genomic DNA may be inserted into BAC vectors which possess both a high copy number origin of replication, which facilitates the isolation of the vector DNA, and a low copy number origin of replication. Cloning of a genomic DNA insert into the high copy number origin of replication inactivates the origin such that clones containing a genomic insert replicate at low copy number. The low copy number of clones having a genomic insert therein permits the inserts to be stably maintained. In addition, selection procedures may be designed which enable low copy number plasmids (i.e. vectors having genomic inserts therein) to be selected. Such vectors and selection procedures are described in the U.S. Patent Application